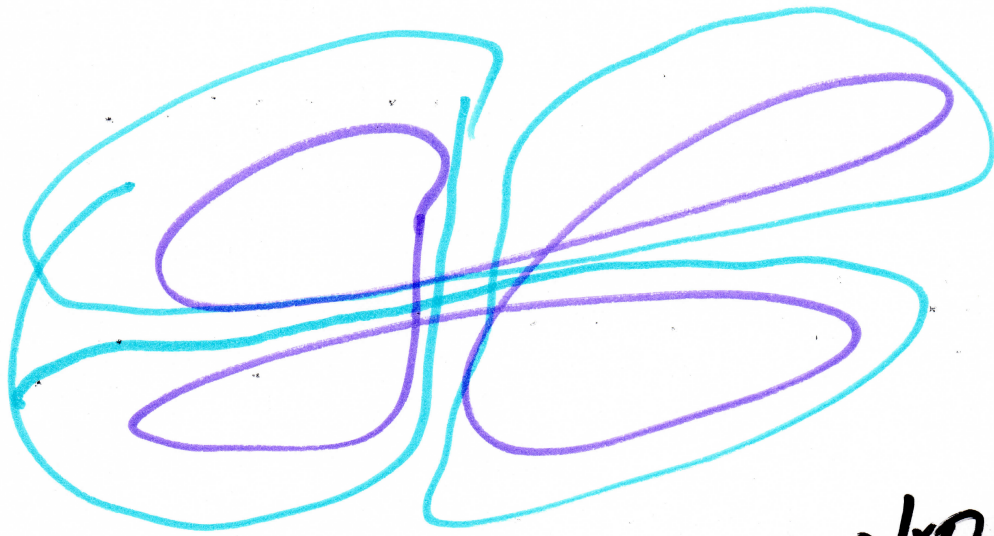


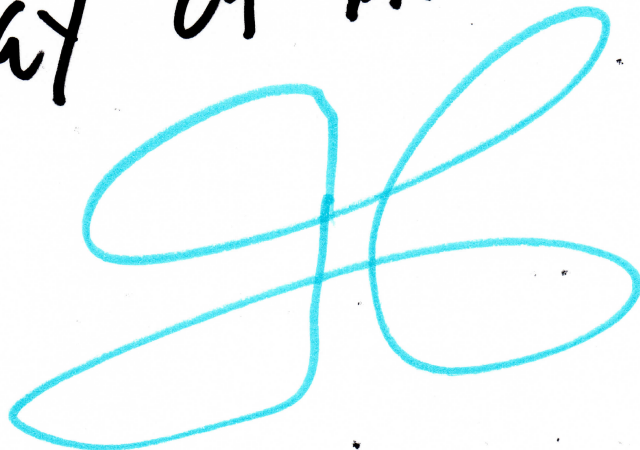
$$\text{Amount of Medicine} = \text{Plasma Concentration} \times \text{Volume of Distribution}$$



by
Takomedi



THE KEY OF PHARMACOKINETICS



2018 NEW

LET'S \$G

PRESCRIBE

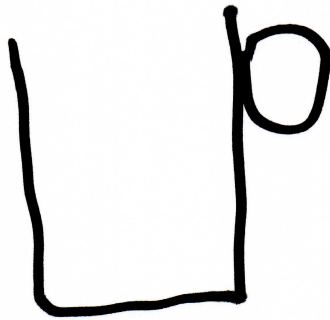
A Thing or a
med

2018

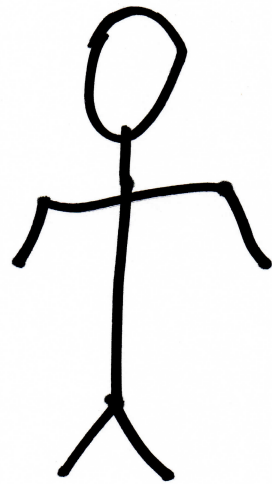
ABDO . B. S. Fatally

① NOV

48 Nov 2018



①



A Container

A Person

Fill
Container
with
water



p ②

A
Person
already
have
a
volume



3



V = volume = Unknown amount

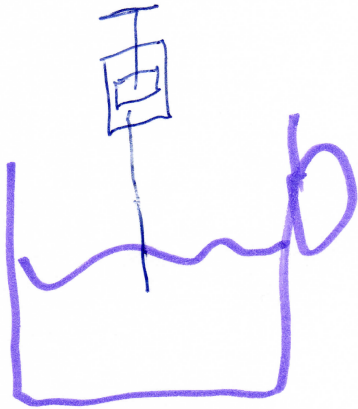
Add
1 gm of
salt
in
 V_1

4

Give
One Tab
of
500mg
Paracetamol

P 3

NOV 2018



C_1



C_2

Draw a sample from
Container C_1 and C_2

C_1 is container

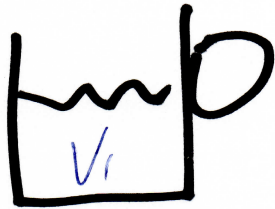
C_2
is
Person

Send both to lab for Test
of
Concentration
D (4)

\$6 Nov

L A B

Results



1mg/ml

8mg
 L

of
Salt
for
1 gm of Salt

of
Paracetamol
for
500 mg

8 Nov 2018

SALT
for 1ml

there is

1mg

50

for

1000mg

there is

1000ml

N

1 LITRE

p6

Paracetamol

for 8mg

there is

1000ml

$V_d =$

Volume of
distribution

is the V_d

51 V_d

50

for

1ml

there is

0.008mg

$$0.008 \times 51 = 0.408$$

$$0.408 \times 1000 = 408$$

408mg of Paracetamol

✱

So given (1) a volume distribution

(2) a plasma concentration

after (1) hour

(3) the parallel mol

concentration is 408

07

✂

To Summarize:

EVERY DRUG HAS a
volume of distribution

Give a known amount

Collect the lab result
for concentration

Multiply $V_d \times \text{concentration}$
THE AMOUNT OF DRUG IN THE
BODY IS THUS OBTAINED ✓

✂

IMPLEMENTATION

When the amount of drug in the body is known at any one time, the toxicity, efficacy, peak level is thus known for maximum treatment against the disease or microorganism

PA

PO

BEGIN

Google a drug
for its

Volume of Distribu-
tion

then take

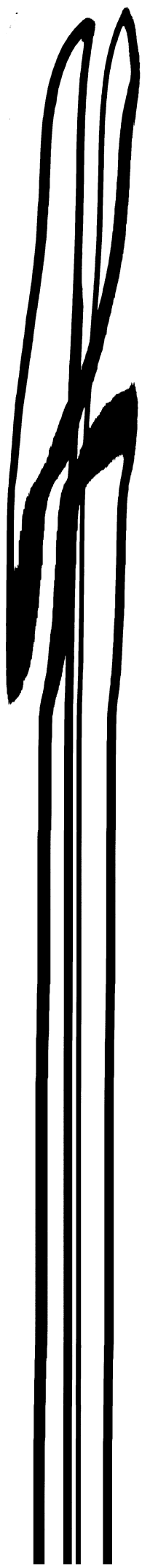
the concentration

and

GET THE Amount of Drug

→

USE THE
Amount of Drug in
the body to
monitor treatment,
toxicity and efficacy



JB

Sunday 4 nov

2018

Sodnal

NMP



Mr. Suran

Drug Absorption

0820
4/11/20

1. Direct diffusion through lipid

2. Pinocytosis

3. Factors

4. BNF

5. Concentration gradient

6. Lipid Solubility

7. Polarity

Non-Polar { No charge

Polar { Charge
positive
negative

8. Polarity & Lipid Solubility

Polar & water soluble

+ , - hydrophilic

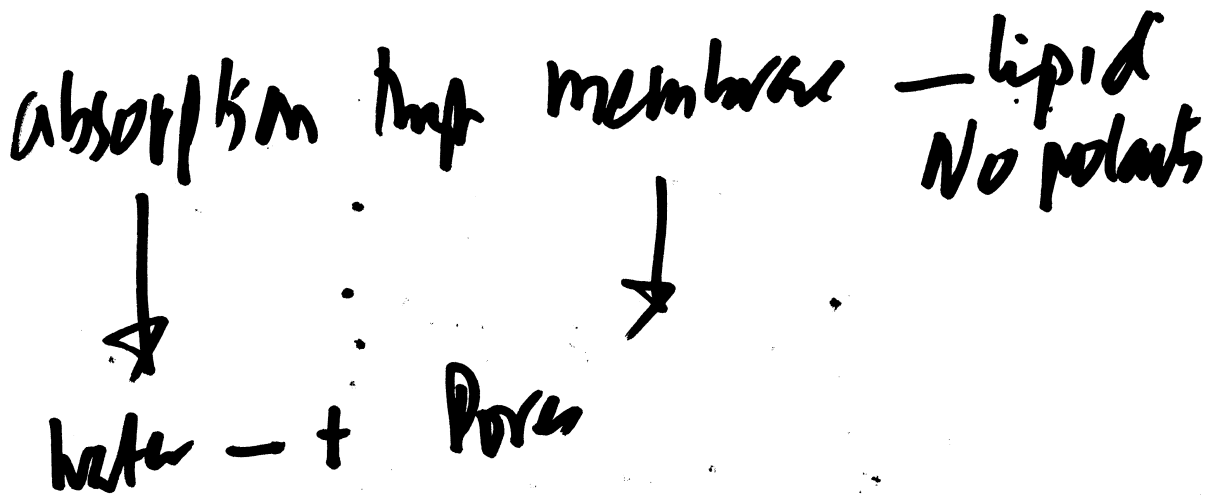
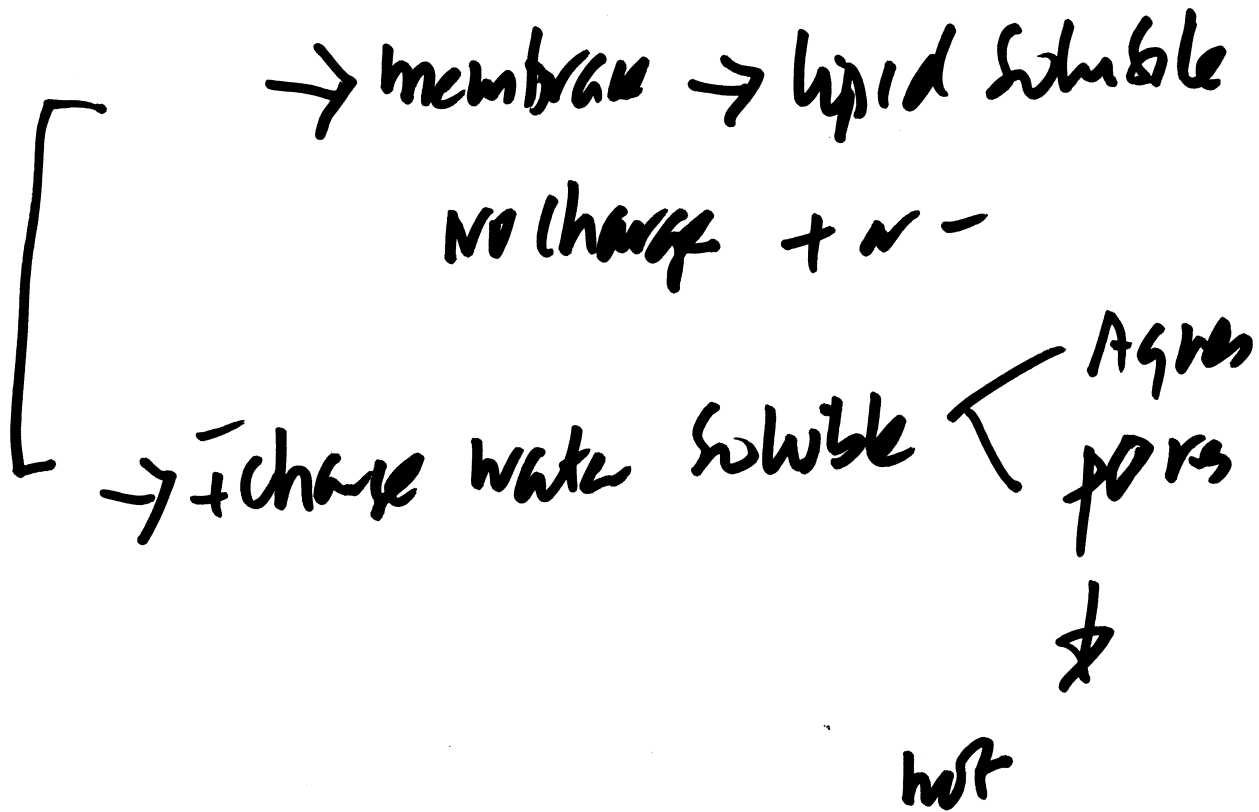
absorbed in H₂O with +, - charge.

①

• KMP

Drug Absorpt

Sören
Linn



②

NAP

IONISATION

Mr Sura

CHAN

+
Lipid soluble

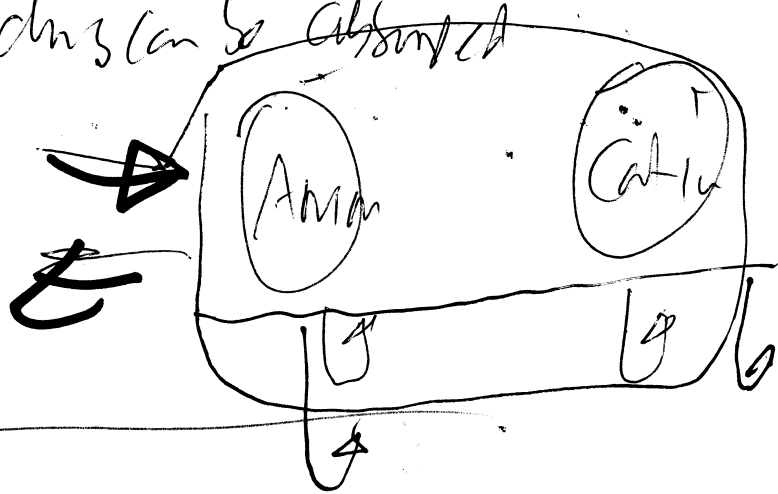
Weak acids - weak base

Ionised = charged

Unionised Uncharged

Only ionized drugs can be absorbed

Unionised
drug



lipid soluble

Ionized forms attract water H_2O

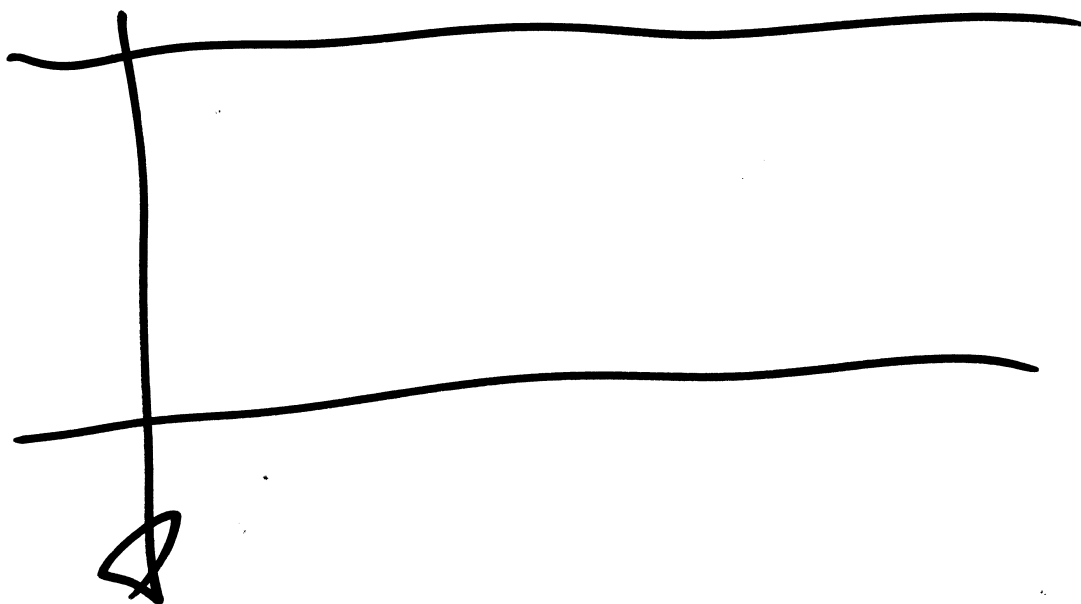
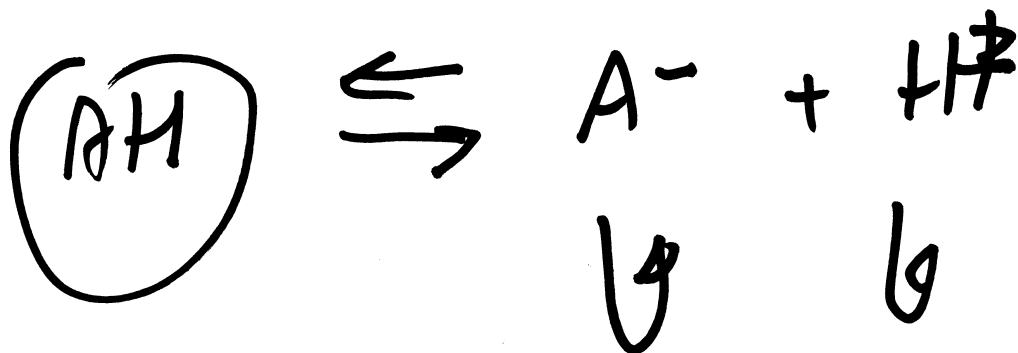
(13)

Sum

88

10/3

Ionization and bond strengths
for a weakly acidic



NHPI

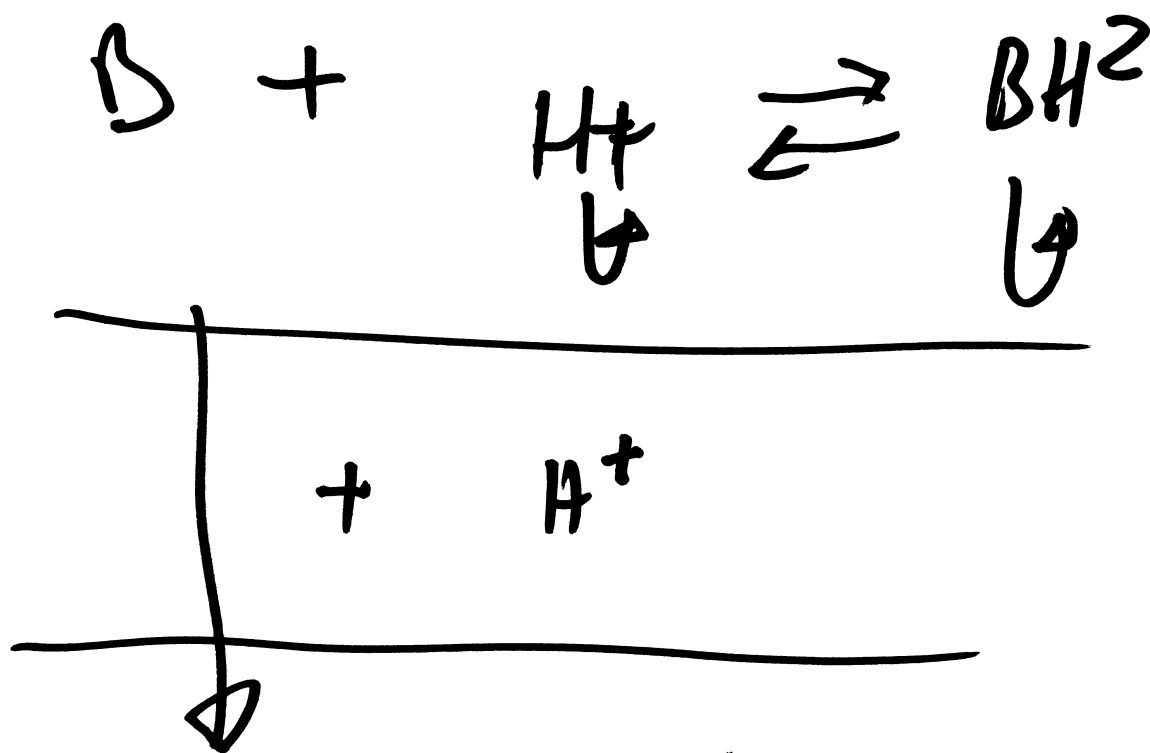
P4

Surv

B

Nov 4

For a weakly basic drug..



NMP

L7 Andic

PG

Effect of pH on Drug GB

Surv

4th

Weakly acid

Aspirin

less ionized

at high pH

Aspirin forms absopts

2 ions determine Acid also P_b ~~Aspirin~~ ~~base~~

Aspirin

Aspirin less ionized low pH

base
Aspirin

less ionized high pH

Aspirin

Aspirin less ionized low pH

Aspirin

Pb

Aspirin

Sur H^+ LAVN
 A solution is all water separation?
 ? ionization is related to acid/base.

} Can an acid become a base

} Can a base become an acid.

if the change in water does not increase.

Acidic less ionized high pH

basic \rightarrow low

Base $<$ ionized \uparrow pH Alkaline

acid less ionized \downarrow

ALD $<$ ionized \downarrow pH Alkaline
 NHD

87

NR4

⊕

⊕

Syrac
m/c

A/B

> <

⊕

H/Ac

Acidic

low

pH

Acid

Base

low

pH

Base
acid

low

pH

A

Acid

B

Alk

8

AMP

is acidic solution base
with ions & charge

GMW

Drug Distribution

Sumen

1. Vascular

2. Extracellular

3. Intracellular

compartment

$$V_d = \frac{\text{Dose}}{\text{concentration in plasma}}$$

multi Equilibria

multi Equilibria

in M Equilibria

$$V_d = \frac{\text{Dose}}{\text{concentration in plasma}}$$

concentration in

plasma.

AMP

PG

hw3

Thurs Disgust.

Suren

The best slide comes to operation a he present that
One shall ahead explain to present this

if he want's 1mg/L from lab

$$\frac{40}{5}$$

$$5L = \frac{10mg}{2mg/L} = 5L \quad 1mg/L \text{ from lab}$$

So sure

$$42L \quad 40L = \frac{5mg}{8mg/L} \quad 40L$$

$$V_d = \frac{10mg}{1mg/L}$$

$$= 10L$$

$$5 \times \frac{8}{1}$$

$$5 \times \frac{1}{8}$$

P10

$$mg T : \frac{mg}{L}$$

$$mg T \times \frac{L}{mg}$$

$$5 \times$$

$$mg : \frac{mg}{L}$$

$$m \times \frac{L}{mg}$$

mg

Dose & Calculations
measured
concentration
1 mg/L
plasma.

Total

$$V_d = \frac{10 \text{ mg}}{1 \text{ mg/L}} = 10 \text{ L}$$

$$1 \text{ mg/L}$$

$$V_d = \frac{D}{C}$$

$$12 \text{ L} = \frac{D}{C}$$

$$= 12 \text{ L} \times 2 \text{ mg/L}$$

$$12 \text{ L} = \frac{D}{0.17 \text{ g/kg}}$$

$$= 12 \text{ L} \times 0.17 \text{ g/kg}$$

P11

$$\frac{L^2}{k} = 12$$

Dosage Calculator

Score
nmv. 24

$$V_d = \frac{\text{Dose}}{\text{mg/L}} = V_d.$$

mg/L



plasma concentration

$$\text{Dose} = V_d \times \text{mg/L}$$

$$\text{mg/L} = \frac{\text{Dose}}{V_d}$$

P12

P

Absorption

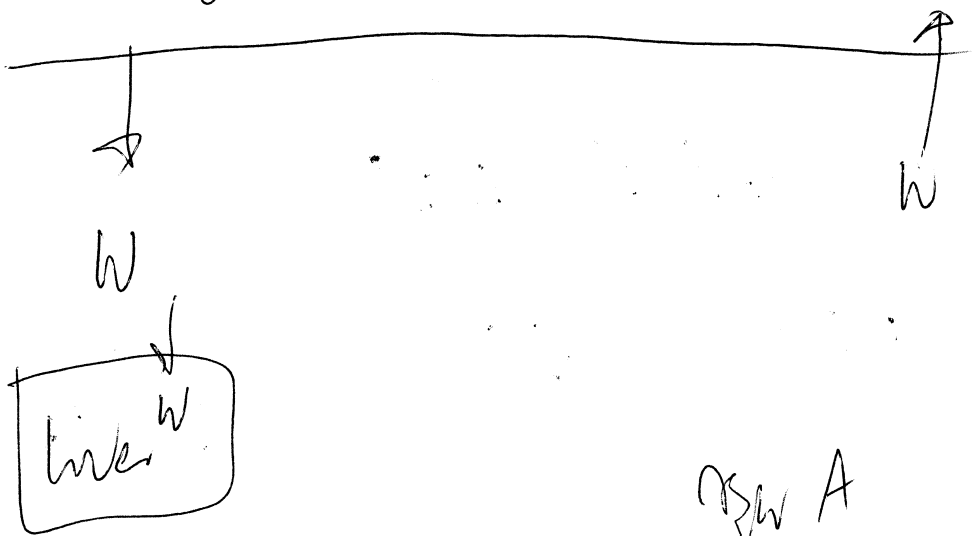
MOV 4

Screen

Plasma protein Binding

Blood

A₀ W



W A

2mL
1mL

Anticoagulant cell

Plasma Protein Binding

Albumin

P13

88

LTC

Nov 4

11:30

Submission of case May, June 2019 was taken -

in his class

1. Supportive services

2. Disease management

3. Case management Holistic

[Under Evaluation of the
history
complaint
circumstances

best in case work
by agreement results
coordinates
114

in problem

row 4

2. Chromine core model

4: The contents of core mode

Why use model #

* mixed model $\begin{matrix} \text{⑩} \\ \swarrow \searrow \\ \text{⑨} \end{matrix} \text{③}$

7 create a model to suit the pts.

15.

76

Mr
Mcken

Long Term Goals List.

Nov 4

- 1 depression
- 2 - alcohol
- 3 drugs
- 4 cigarettes
- 5 Diabetes
- 6 - Cancer
- 7 -

LTC

16